taking into account the neighboring Fe-atoms up to the 6th nearest. The calculated value was estimated as 4.0×10^5 erg/cm³ at 0 K. The temperature dependence of K_1 taken into account K_d do not obey the third power function supposed for the crystalline field anisotropy. The value of K_1 may be due to the anisotropic exchange as well as the crystalline field anisotropy.

Since the ferromagnetic-paramagnetic transition at T_c is accompanied by the first order transition with a lattice distortion, the pressure variation of T_c , $\Delta T_c/\Delta p$, has to satisfy the thermodynamical Clausius Clapeyron equation;

$$\frac{\Delta T_{\rm c}}{\Delta p} = \frac{\Delta V}{\Delta S},\tag{3}$$

where ΔS and ΔV are the discontinuous jumps in the entropy and the volume at $T_{\rm e}$, respectively. We have measured the variation of $T_{\rm e}$ under the hydrostatic pressures up to 6 kbar and obtained as $\Delta T_{\rm e}/\Delta P = -3.46 \times 10^{-3}$ deg/ bar. Using the value of ΔV obtained in this experiment, the magnitude of ΔS is estimated as 0.04 cal/mol·deg, which is relatively small for a first-order transition.

According to Rodbell and Bean,¹⁷⁾ if the exchange interaction which give rise to the magnetic ordering strongly depends upon the interatomic specing, the Curie temperature T_c is expressed by the following equation,

$$T_{\rm c} = T_0 \left(1 + \beta \, \frac{V - V_0}{V_0} \right), \tag{4}$$

where V_0 is the specific volume in the absence of exchange interaction, T_0 the Curie temperature of the rigid lattice and β the ratio between the changes of T_c and volume. On the basis of the molecular field approximation, the exchange striction at 0 K from the equilibrium condition is given by

$$\left(\frac{V-V_0}{V}\right) = \frac{3}{2} \frac{j^2}{j(j+1)} NkKT_0\beta.$$
 (5)

Where j is the total angular momentum, K is the compressibility, N is the number of magnetic atoms per unit volume, k is the Boltzman constant. Here, if we define the parameter as

$$\eta = 40NkKT_0\beta^2[j^2(j+1)^2/(2j+1)^4 - 1], \quad (6)$$

we notice that for $\eta < 1$ the transition is of second order, while for $\eta > 1$, the transition is

of first order. If the ferromagnetic-paramagnetic transition is of first order, the volume change at T_c is given by

$$\left(\frac{\Delta V}{V}\right) = \frac{3}{2} \frac{j^2}{j(j+1)} Nk K T_0 \beta \sigma_c^2, \qquad (7)$$

where σ_c is the jump of the relative magnetization at T_c . Also, the pressure variation of the Curie temperature can be derived from eq. (4) as,

$$\frac{\partial T_{\rm c}}{\partial p} = -\beta K T_0. \tag{8}$$

We can estimate the exchange striction at 0 K and the volume change at T_c from eqs. (5), (6), (7) and (8). On substituting the value of K, T_0 , and $\Delta T_c/\Delta P$ into eq. (8), we have the value of $\beta = 28.8$. Here, we used the value of $K = 4.8 \times$ 10^{-13} cm²/dyne for Fe₂P measured by means of the X-ray diffraction under high pressure, which will be reported in near future, the value of $T_0 = 250$ K obtained by extrapolating the Brillouin function curve for $j = \frac{3}{2}$ to zero magnetization. On substituting the values of $\beta =$ 28.8 and $j = \frac{3}{2}$ deduced by the magnetization at 0 K in eq. (6) we have $\eta = 1.20$. This suggests that the transition at T_c is of first order. The exchange striction at 0 K and the jump of volume at T_c are estimated to be 9.1×10^{-3} and 1.6×10^{-3} , from eqs. (5) and (7), respectively. Here, the value of $\sigma_{\rm e}$ was used 0.44 obtained from the jump of the hyperfine field at $T_{\rm c}$ determined by Wäppling et al.11) These estimated values are in reasonable agreement with the experimental values of 9.5×10^{-3} and 0.64×10^{-3} , respectively. In conclusion, the ferromagnetic to paramagnetic transition at $T_{\rm c}$ of Fe₂P might be accompanied by the firstorder transition with a lattice distortion due to magnetoelastic effects.

References

- S. Rundquist and F. Jellinek: Acta Chem. Second. 13 (1956) 425.
- B. Carlsson, M. Gölin and S. Rundqvist: J. Solid State Chem. 8 (1973) 57.
- 3) S. Chiba: J. Phys. Soc. Japan 15 (1960) 581.
- A. J. P. Meyer and M. C. Cadeville: J. Phys. Soc. Japan 17 (1962) 223.
- R. Fruchart, A. Roger and J. P. Senateur: J. appl. Phys. 40 (1969) 1250.
- D. Bellavance, J. Mikkelsen and A. Wold: J. Solid State Chem. 2 (1970) 285.
- 7) R. Wäppling, L. Häggystrom, S. Rundqvist and

1977)

E. Karlsson: J. Solid State Chem. 3 (1971) 276.

- L. Lundgren, C. Beckman, G. Tarmohammed and S. Rundqvist: 4th Int. Conf. Solid Compounds Transition Elements (1973) p. 68.
- K. Sato, K. Adachi and E. Ando: J. Phys. Soc. Japan 26 (1969) 855.
- 10) R. E. Bailey and J. F. Duncan: Inorg. Chem. 6 (1966) 1444.
- R. Wäppling, L. Häggstrom, T. Ericsson, S. Devanarayanan, E. Karlsson, B. Carlsson and S. Rundqvist: J. Solid State Chem. 13 (1975) 258.
- C. Inoue, H. Kadomatsu, H. Fujii and T. Okamoto: J. Phys. Soc. Japan 35 (1973) 1322.
- 13) J. B. Goodenough: J. Solid State Chem. 7 (1973) 428.
- 14) N. P. Grazhadankina, E. A. Zavadshii and I. G. Fakidov: Fiz. tverdogo Tela 11 (1969) 2327.
- S. Nagase, H. Watanabe and T. Shinohara: J. Phys. Soc. Japan 34 (1973) 908.
- 16) C. Zener: Phys. Rev. 96 (1954) 1355.
- 17) D. S. Rodbell and C. P. Bean: J. appl. Phys. 33 (1962) 1037.

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